Amendments to the Claims:

This listing of claims will replace all prior versions, and listings, of claims in the application:

Listing of Claims:

Claims 1-13 (cancelled)

Claim 14 (currently amended): A compound which is 1-[2-(2-ethyl-2H-tetrazol-5-yl)-ethyl]-3-[5-(3-fluoro-4-methanesulfonyl-phenyl)-4-methyl-thiazol-2-yl]-urea of formula I

in free or salt form, wherein

 R^a -is hydrogen or C_1 - C_4 -alkyl, R^b -is C_1 - C_8 -alkyl substituted by pyridyl, R^3 -is R^6 , and R^4 -is fluoro or C_1 - C_8 -haloalkyl,

or R^a is hydrogen or C_1 - C_4 -alkyl, R^b is C_1 - C_8 -alkyl substituted by hydroxy or nitrile, R^3 -is R^6 , and R^4 is hydrogen or C_1 - C_8 -haloalkyl,

or R^a -is hydrogen or C_1 - C_4 -alkyl, R^b -is C_1 - C_8 -alkyl substituted by nitrile, R^3 -is fluoro, and R^4 -is R^7 ,

or R^a is hydrogen or C_1 - C_4 -alkyl, R^b is C_1 - C_8 -alkyl-substituted by hydroxy, R^3 is fluoro, and R^4 is R^7 ,

or R^a is hydrogen or C₁-C₄-alkyl, R^b is C₁-C₈-alkyl substituted by di(C₁-C₈-alkyl)amino, R³ is R⁶, and R⁴ is C₁-C₈-haloalkyl,

or R^a is hydrogen or C₁-C₄-alkyl, R^b is C₁-C₈-alkyl substituted by O-C₁-C₈-alkyl-OH, R³ is R⁶, and R⁴ is fluoro or C₁-C₈-haloalkyl,

or R^a is hydrogen or C₁-C₄-alkyl, R^b is -CH(CH₃)-CH₂-OH, R³ is R⁶, and R⁴ is fluoro,

or R^a is hydrogen or C₁-C₄-alkyl, R^b is C₁-C₈-alkyl substituted by pyrrolidinyl substituted by C₁-C₈-alkyl, R³ is R⁶, and R⁴ is C₁-C₈-haloalkyl,

or R^a is hydrogen or C_1 - C_4 -alkyl, R^b is C_1 - C_8 -alkyl substituted by oxazolyl substituted by C_1 - C_8 -alkyl, R^3 is R^6 , and R^4 is nitrile or imidazolyl, or R^a is hydrogen or C_1 - C_4 -alkyl, R^b is C_1 - C_8 -alkyl substituted by imidazolyl, R^3 is R^6 , and R^4 -is fluoro,

or R^a is hydrogen or C_1 - C_4 -alkyl, R^b is C_1 - C_8 -alkyl substituted by benzoimidazolyl, R^3 is R^6 , and R^4 is fluoro,

or R^a is hydrogen or C_1 - C_4 -alkyl, R^b is C_1 - C_8 -alkyl substituted by isoxazolyl substituted by C_1 - C_8 -alkyl, R^3 is R^6 , and R^4 is R^7 ,

or R^a is hydrogen or C_1 - C_4 -alkyl, R^b is C_1 - C_8 -alkyl substituted by pyrrolyl substituted by C_1 - C_8 -alkyl, R^3 is R^6 , and R^4 is R^7 ,

or R^a -is hydrogen or C_1 - C_4 -alkyl, R^b -is C_1 - C_8 -alkyl substituted by pyrazolyl substituted by C_1 - C_8 -alkyl, R^3 -is R^6 -and R^4 -is R^7 .

or R^a is hydrogen or C_1 - C_4 -alkyl, R^b is C_1 - C_8 -alkyl substituted by -CO-O-CH₃, -CO-O-butyl, -CO-di(C_1 - C_8 -alkyl)amino, -CO-NH₂, -NH-CO- C_1 - C_8 -alkyl, -SO₂- C_1 - C_8 -alkyl, -CO-NH- R^6 where R^6 is napthyl, or by -CO-NH- C_1 - C_8 -alkyl optionally substituted by di(C_1 - C_8 -alkyl) amino, R^3 is R^6 , and R^4 is R^7 .

or R^a -is hydrogen or C_1 - C_4 -alkyl, R^b -is -CH(CH₃)-CO-NH- C_1 - C_8 -alkyl or -CH(CH₃)-CO-O- C_1 - C_8 -alkyl, R^3 -is- R^6 , and R^4 -is R^7 ,

or R^a is hydrogen or C_1 - C_4 -alkyl, R^b -is C_1 - C_8 -alkyl substituted by -CH(OH)-CH₂-OH, R^3 is R^6 , and R^4 is R^7 ,

or R^a is hydrogen or C_1 - C_4 -alkyl, R^b is C_1 - C_8 -alkyl substituted by C_1 - C_8 -alkoxy, or by -S- C_1 - C_8 -alkyl, R^3 is R^6 , and R^4 is R^7 ,

or R^a is hydrogen or C₁-C₄-alkyl, R^b is C₁-C₈-alkyl substituted by a 5- or 6-membered heterocyclic ring having one or more ring hetero atoms selected from the group consisting of oxygen, nitrogen and sulphur, that ring being substituted by oxo, R³ is R⁶, and R⁴ is R⁷,

or R^a is hydrogen or C₁-C₄-alkyl, R^b is C₁-C₈-alkyl substituted by a 5- or 6-membered heterocyclic ring having three or more ring hetero atoms selected from the group consisting of oxygen, nitrogen and sulphur, that ring being optionally substituted by C₁-C₈-alkyl, -C₁-C₈-alkyl-di(C₁-C₈-alkyl)amino, or by C₂-C₈-cycloalkyl, R³ is R⁶, and R⁴ is R⁷,

or R^a is hydrogen or C_1 - C_4 -alkyl, R^b is C_1 - C_8 -alkyl substituted by oxazolyl substituted by C_3 - C_8 -alkyl, R^3 is R^6 , and R^4 is R^7 ,

or R^a is hydrogen or C_1 - C_4 -alkyl, R^b is C_1 - C_8 -alkyl substituted by imidazolyl substituted by C_1 - C_8 -alkyl optionally substituted by hydroxy or C_1 - C_8 -alkoxy, R^3 is R^6 , and R^4 is R^7 ,

or R^a-is hydrogen or C₁-C₄-alkyl, R^b-is C₁-C₈-alkyl substituted by -CO-Het where Het is a 5-or 6-membered heterocyclic ring having two or more ring hetero atoms selected from the group consisting of oxygen, nitrogen and sulphur, that ring being optionally substituted by C₁-C₈-alkyl, R³ is R⁶, and R⁴ is R⁷.

or R^a is hydrogen or C₁-C₄-alkyl, R^b is a 5- or 6-membered heterocyclic ring having one or more ring hetero atoms selected from the group consisting of oxygen, nitrogen and sulphur, that ring being substituted by oxo, R³ is R⁶, and R⁴ is R⁷.

or R^a is hydrogen or C₁-C₄-alkyl, R^b is an aza-bicyclo[3.2.1]oct-3-yl ring optionally substituted by C₁-C₈-alkyl, R³ is R⁶, and R⁴ is R⁷,

or R^a and R^b together form an azetidine ring substituted by C_4 - C_8 -alkoxycarbonyl or nitrile, R^3 is R^6 , and R^4 is R^7 .

or R^a and R^b together form a pyrrolidine ring substituted by -CO-NH₂ or nitrile, R³ is R⁶, and R⁴ is R⁷,

or R^a and R^b together form an imidazo-pyridine ring, R³ is R⁶, and R⁴ is R⁷;

R² is C₁-C₄-alkyl or halogen;

R⁵-is hydrogen, halogen or C₁-C₈-alkyl;

R⁶-is halo, -SO₂-CH₃, -SO₂-CF₃, carboxy, -CO-NH₂, -CO-di(C₁-C₈-alkyl)amino, or a 5- or 6-membered heterocyclic ring having one or more ring hetero atoms selected from the group consisting of oxygen, nitrogen and sulphur, that ring being optionally substituted by halo, cyano, oxo, hydroxy, carboxy, nitro, C₃-C₈-cycloalkyl, C₁-C₈-alkylcarbonyl, C₁-C₈-alkoxy optionally substituted by aminocarbonyl, or C₁-C₈-alkyl optionally substituted by hydroxy, C₁-C₈-alkoxy, C₁-C₈-alkylamino;

 R^7 is hydrogen, halo, $-SO_2$ -CH₃, nitrile, C_1 -C₈-haloalkyl, imidazolyl, C_1 -C₈-alkyl, $-NR^8R^9$, or $-SO_2$ -NR⁸R⁹; and

R⁸-and R⁹-are independently hydrogen, amino, C₁-C₈-alkylamino, di(C₁-C₈-alkyl)amino, or C₁-C₈-alkyl optionally substituted by hydroxy, or R⁸-and R⁹ together form a 5- to 10-membered heterocyclic ring having one or more ring hetero atoms selected from the group consisting of oxygen, nitrogen and sulphur, that ring being optionally substituted by halo, cyano, oxo, hydroxy, carboxy, nitro, C₃-C₈-cycloalkyl, C₁-C₈-alkyloanino, oxogen, oxogen, nitrogen and sulphur, that ring being optionally substituted by halo, cyano, oxogen, hydroxy, nitro, C₃-C₈-cycloalkyl, C₁-C₈-alkyloanino, oxogen, nitrogen and sulphur, that ring being optionally substituted by halo, cyano, oxogen, hydroxy, nitro, C₃-C₈-cycloalkyl, C₁-C₈-alkyloanino, oxogen, nitrogen and sulphur, that ring being optionally substituted by halo, cyano, oxogen, hydroxy, nitro, C₃-C₈-cycloalkyl, C₁-C₈-alkyloanino, oxogen, nitrogen and sulphur, that ring being optionally substituted by halo, cyano, oxogen, hydroxy, nitro, C₃-C₈-cycloalkyl, C₁-C₈-alkyloanino, oxogen, nitrogen and sulphur, that ring being optionally substituted by hydroxy, C₁-C₈-alkoxy, C₁-C₈-alkyloanino oxogen, nitrogen and sulphur, that ring being optionally substituted by aminocarbonyl, oxogen, nitrogen and sulphur, that ring being optionally substituted by aminocarbonyl, oxogen, nitrogen and sulphur, that ring being optionally substituted by aminocarbonyl, oxogen, nitrogen and sulphur, that ring being optionally substituted by aminocarbonyl, oxogen, nitrogen and sulphur, that ring being optionally substituted by aminocarbonyl, oxogen, nitrogen and sulphur, that ring being optionally substituted by aminocarbonyl, oxogen, nitrogen and sulphur, that ring being optionally substituted by aminocarbonyl, oxogen, nitrogen and sulphur, that ring being oxogen, nitrogen and sulphur, that ring being oxogen, nitrogen and sulphur, that ring being oxogen, nitrogen and nitrogen and nitrogen and nitrogen and nitrogen and nitrogen and nitrogen and

Claims 15-19 (cancelled)

Claim 20 (currently amended): A pharmaceutical composition comprising a compound according to claim 14, in free or salt form.

Claim 21 (withdrawn-currently amended): A method of treating a disease mediated by phosphatidylinositol 3-kinase in a subject in need of such treatment, which comprises administering to said subject an effective amount of a compound of formula I as defined in claim 14 in free form or in the form of a pharmaceutically acceptable salt.

Claim 22 (withdrawn-currently amended): A method of treating respiratory diseases, allergies, rheumatoid arthritis, osteoarthritis, rheumatic disorders, psoriasis, ulcerative colitis, Crohn's disease, septic shock, proliferative disorders such as cancer, atherosclerosis, allograft rejection following transplantation, diabetes, stroke, obesity or restenosis in a subject in need of such treatment, which comprises administering to said subject an effective amount of a compound of formula I as defined in claim 14 in free form or in the form of a pharmaceutically acceptable salt.

Claim 23 (withdrawn-currently amended): A process for the preparation of preparing a compound of formula I as defined in claim 14, in free or salt form which comprises the steps of:

(i) (A) reacting a compound of formula II

$$\mathbb{R}^{3}$$
 \mathbb{R}^{4}
 \mathbb{R}^{4}
 \mathbb{R}^{5}
 \mathbb{R}^{4}
 \mathbb{R}^{5}
 \mathbb{R}^{4}
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 \mathbb{R}^{4}
 \mathbb{R}^{4}

wherein R^2 , R^3 , R^4 and R^5 are as claimed in claim 14 and T is a 5- or 6-membered heterocyclic ring having one or more ring hetero atoms selected from the group consisting of oxygen, nitrogen and sulphur, with a compound of formula III

wherein R^a and R^b are as claimed in claim 14; or

(B) reacting eompounds a compound of formula IV

wherein R², R³, R⁴ and R⁵ are as claimed in claim 14 with a compound of formula III
wherein R^a and R^b are as claimed in claim 14; or

(C) for the preparation of compounds of formula I where R^a is hydrogen and R^2 , R^3 , R^4 , R^5 and R^b are as claimed in claim 14, reacting a compound of formula V

wherein R², R³, R⁴ and R⁵ are as claimed in claim 14, with a compound of formula VI

wherein R^b is as claimed in claim 14; or

(D) for the preparation of compounds of formula I where R^a -is-hydrogen, R^b -is C_1 - C_8 -alkyl substituted by imidazolyl substituted by C_1 - C_8 -alkyl optionally substituted by hydroxy or C_1 - C_8 -alkoxy and R^2 , R^3 , R^4 -and R^5 -are as claimed in claim 14, reacting a compound of formula V where R^2 , R^3 , R^4 and R^5 are as claimed in claim 14, with a compound of formula VII

where Q is C₁-C₈-alkyl optionally substituted by hydroxy or C₁-C₈-alkoxy; and

(ii) recovering the resultant compound of formula I in free or salt form.